

Inherited variants affecting RNA editing may contribute to ovarian cancer susceptibility: results from a large-scale collaboration

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ABSTRACT

RNA editing in mammals is a form of post-transcriptional modification in which adenosine is converted to inosine by the adenosine deaminases acting on RNA (ADAR) family of enzymes. Based on evidence of altered ADAR expression in epithelial ovarian cancers (EOC), we hypothesized that single nucleotide polymorphisms (SNPs) in ADAR genes modify EOC susceptibility, potentially by altering ovarian tissue gene expression. Using directly genotyped and imputed data from 10,891 invasive EOC cases and 21,693 controls, we evaluated the associations of 5,303 SNPs in *ADAD1*, *ADAR*, *ADAR2*, *ADAR3*, and *SND1*. Unconditional logistic regression was used to estimate odds ratios (OR) and 95% confidence intervals (CI), with adjustment for European ancestry. We conducted gene-level analyses using the Admixture Maximum Likelihood (AML) test and the Sequence-Kernel Association test for common and rare variants (SKAT-CR). Association analysis revealed top risk-associated SNP rs77027562 (OR (95% CI)= 1.39 (1.17-1.64), $P=1.0 \times 10^{-4}$) in *ADAR3* and rs185455523 in *SND1* (OR (95% CI)= 0.68 (0.56-0.83), $P=2.0 \times 10^{-4}$). When restricting to serous

histology ($n=6,500$), the magnitude of association strengthened for rs185455523 ($OR=0.60$, $P=1.0 \times 10^{-4}$). Gene-level analyses revealed that variation in *ADAR* was associated ($P<0.05$) with EOC susceptibility, with $P_{AML}=0.022$ and $P_{SKAT-CR}=0.020$. Expression quantitative trait locus analysis in EOC tissue revealed significant associations ($P<0.05$) with *ADAR* expression for several SNPs in *ADAR*, including rs1127313 (G/A), a SNP in the 3' untranslated region. In summary, germline variation involving RNA editing genes may influence EOC susceptibility, warranting further investigation of inherited and acquired alterations affecting RNA editing.

INTRODUCTION

Over the past decade it has been recognized that the complexity of higher organisms is related to the information stored in non-protein-coding regions of the genome. Such complexity may be attributed to a range of processing events and post-transcriptional modifications that affect the fate of RNA, including alternative splicing, 5' capping, 3' polyadenylation, and RNA editing [1-3]. The most common type of RNA editing in eukaryotes is site-selective hydrolytic deamination of adenosine into inosine (A-to-I) within double-stranded RNAs, and recent bioinformatic analyses and high-throughput sequencing efforts have revealed that A-to-I editing is widespread and alters non-coding and protein-coding sequences throughout the genome [4].

A-to-I editing is mediated by a family of adenosine deaminases acting on RNA (ADARs), and this process modulates expression of genes and biological pathways *via* several mechanisms [4]. Indeed, altered expression and/or activity of ADAR enzymes has been linked to a variety of conditions, including cardiovascular and neurological diseases and cancers [4]. Epithelial ovarian cancer (EOC) is the fifth leading cause of cancer death among women in the United States [5], and ADAR expression levels have been reported to be significantly higher in serum and peritoneal fluid from patients with EOCs compared with benign ovarian tumors [6, 7], suggesting ADARs may be useful biomarkers for the diagnosis and management of EOC.

We hypothesized that germline single nucleotide polymorphisms (SNPs) involving ADAR-related/RNA editing genes may contribute to EOC risk. The main purpose of this investigation was to determine whether SNPs in five ADAR genes (*ADAD1*, *ADAR*, *ADAR2*, *ADAR3*, and *SND1*) were associated with EOC susceptibility. We used data available from a large-scale genotyping collaboration involving 10,891 EOC cases and 21,693 controls from the international Ovarian Cancer Association Consortium (OCAC) [8]. We also sought to evaluate the overall contribution of each gene on EOC susceptibility and to determine whether candidate SNPs associated with altered expression of corresponding genes in EOC tumor tissue.

RESULTS

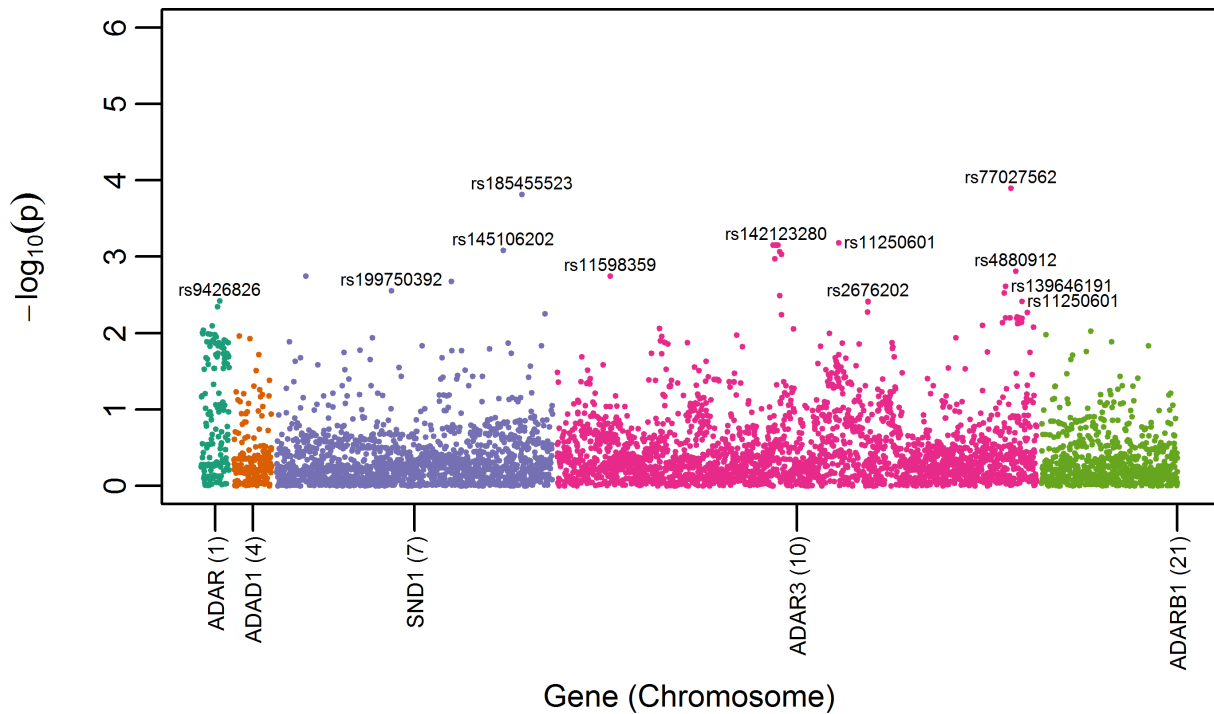
Study population

The study sample included 10,891 invasive EOC patients and 21,693 controls of European ancestry (Supplementary Table 1). Selected subject characteristics are shown in Table 1. The mean age at diagnosis for cases was 58.1 years, the mean age at interview for controls was 56.1 years. Cases were more likely than controls to be nulliparous and to have never used oral contraceptives. Most cases had serous histology (59.7%), distant stage (63.0%), and high-grade disease (58.9%).

Variant-level association analysis and overlap with regulatory domains

SNP-level association analysis revealed top-ranked SNPs (defined as the top 5% of SNPs having the most statistically significant P values) in *ADAR*, *ADAR3*, and *SND1* in the all-histologies and serous-only analyses (Figure 1A and 1B). Table 2 summarizes association results for the most statistically significant SNPs overall or by serous histology ($P < 4.0 \times 10^{-3}$); associations were not significant after correction for multiple testing ($FDR > 0.15$). Most of the top-ranked variants were imputed, rare or low frequency ($MAF < 0.05$), and not part of a shared haplotype. rs77027562 (A>G; $MAF = 0.009$), the top risk-associated variant among all histologies (OR (95% CI) = 1.39 (1.17-1.64), $P = 1.0 \times 10^{-4}$), resides in an intron of *ADAR3*. *ADAR3* SNP rs77027562 and its proxies ($r^2 > 0.80$) reside in genomic regions that overlap with regulatory domains, particularly enhancers in blood and brain (Table 3). The next top-ranked variant, *SND1* rs185455523 (T>A), was associated with a decreased EOC risk (OR (95% CI) = 0.68 (0.56-0.83), $P = 1.5 \times 10^{-4}$), but this SNP and its proxies do not appear to overlap with regulatory domains. When analysis was restricted to the 6,500 patients with invasive serous adenocarcinomas, the magnitude of association was slightly attenuated for *ADAR3* rs77027562 ($OR = 1.33$, $P = 6.1 \times 10^{-3}$) and slightly stronger for *SND1* rs185455523 ($OR = 0.60$, $P = 1 \times 10^{-4}$). Exploratory analysis for the less common histologic subtypes (endometrioid ($n = 1,439$), mucinous

A) All Invasive



B) Serous

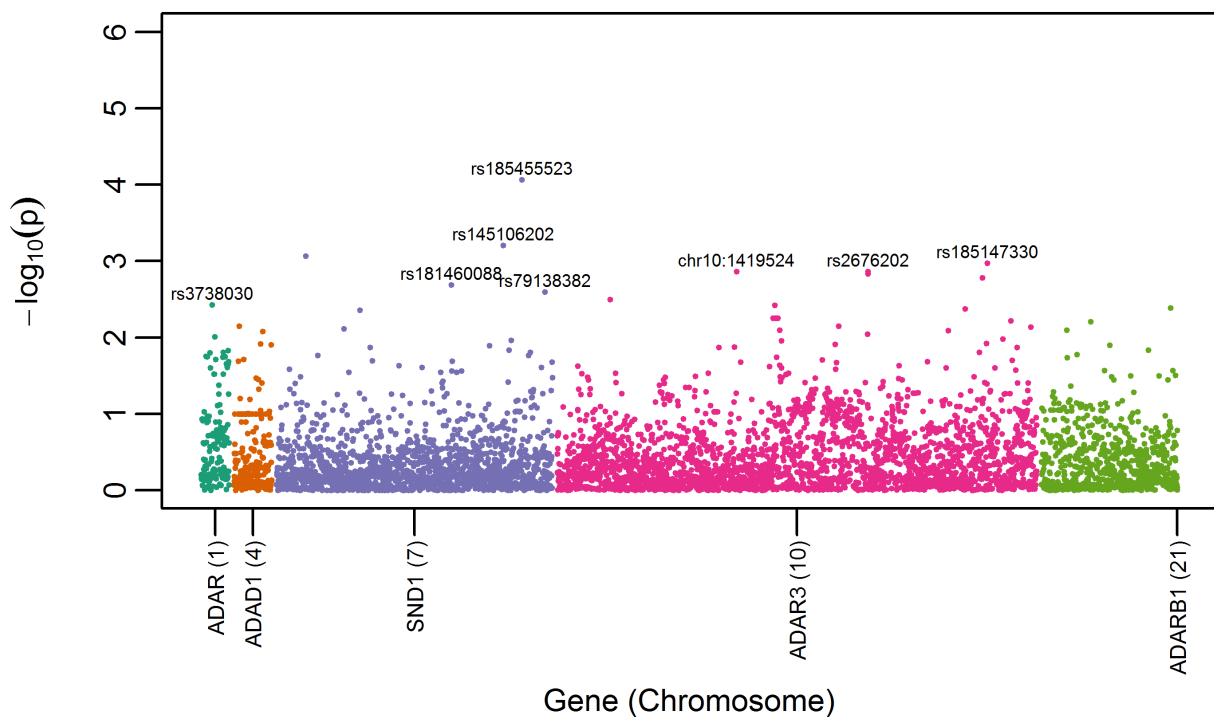


Figure 1: Manhattan plot for candidate RNA editing SNPs among a) all invasive cases ($n = 10,891$) versus controls ($n = 21,693$) and b) serous cases ($n = 6,500$) versus controls.

Table 1: Characteristics of study participants (N = 32,584)

Variable	Cases (n = 10,891)	Controls (n = 21,693)
Age at diagnosis/interview(y), mean (SD)	58.1 (11.4)	56.1 (24.9)
History of pregnancy		
Yes	6021 (80.4)	15190 (87.9)
No	1318 (17.6)	1868 (10.8)
Unknown	149 (2.0)	217 (1.3)
Oral contraceptive use		
Ever	4017 (57.4)	10572 (63.3)
Never	2864 (41.0)	5900 (35.3)
Unknown	112 (1.6)	243 (1.5)
Histology		
Serous	6500 (59.7)	NA
Mucinous	696 (6.4)	
Endometrioid	1439 (13.2)	
Clear Cell	660 (6.1)	
Mixed Cell	369 (3.4)	
Other or unknown epithelial type	1227(11.3)	
Stage		
Localized	1425 (15.7)	NA
Regional	1838 (20.2)	
Distant	5721 (63.0)	
Unknown	103 (1.1)	
Grade		
I/II	2882 (32.8)	NA
III/IV	5174 (58.9)	
Other/Unknown	729 (8.3)	

(n = 696), and clear cell (n = 660)) revealed several SNP-level associations unique to each sub-type (Figure 2A-2C). For example, rs145678553-C in *ADAR3* is a rare variant (MAF = 0.0047) associated with an increased risk for mucinous EOC (OR (95%CI) = 3.46 (1.91-6.26), $P = 3.99 \times 10^{-5}$), and rs116983191-A in *ADAR3* is a low-frequency variant (MAF = 0.044) associated with clear cell carcinoma (OR (95%CI) = 1.86 (1.42-2.43, $P = 6.91 \times 10^{-6}$). rs145678553-C was not represented in Haploreg. rs116983191-A is located in promoter and enhancer regions, but not in tissues relevant to ovarian cancer.

Gene-level analyses

Gene-level analyses based on AML and SKAT-CR revealed that variation in *ADAR* was nominally associated ($P < 0.05$) with susceptibility to all invasive EOC, with $P = 0.02$ using both methods (Table 4). Histology-specific analyses revealed that *ADAR* variation was associated with endometrioid EOC susceptibility ($P_{\text{SKAT-CR}} = 0.005/P_{\text{AML}} = 0.008$). When using a Bonferroni threshold of 0.0025, only *ADAR3* variation was significantly associated with mucinous histology ($P_{\text{SKAT-CR}} = 0.0016/P_{\text{AML}} = 0.031$).

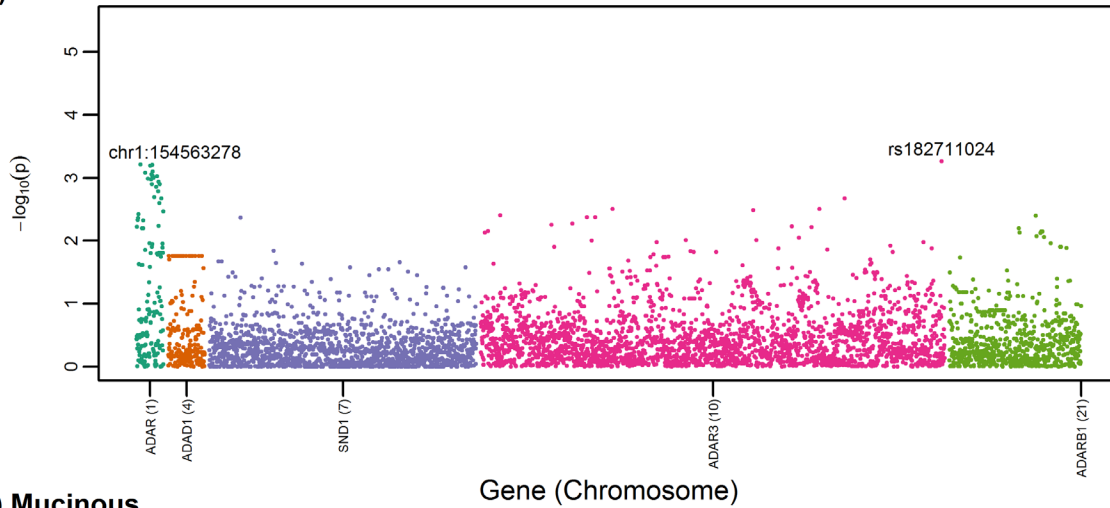
To examine associations between genotype and gene expression for the 5 candidate RNA editing genes, expression quantitative trait locus (eQTL) analysis was performed using matched genotype and tissue expression data from The Cancer Genome Atlas (TCGA) high-grade

serous adenocarcinoma tumors (<https://tcga-data.nci.nih.gov/tcga/>). eQTL analysis revealed statistically significant associations ($P < 0.05$) with *ADAR* expression for several SNPs in *ADAR*, including rs1127313 (G/A), a SNP in the 3'UTR within a putative miRNA binding site that was associated with susceptibility in all histologies (OR = 1.05, $P = 0.009$). rs1127313 is also in high LD ($r^2 = 0.86$) with top *ADAR* risk SNP rs9426826 (see Table 2). *ADAR* tumor tissue expression was slightly higher among G allele carriers of rs1127313 compared to A allele carriers ($P = 0.027$; Figure 3). rs1127313 is also an eQTL for *ADAR* in whole blood (Supplementary Table 2), and lies in a genomic region with enhancer features and DNase I hypersensitivity site in several tissues, including ovary. Statistically-significant cis-eQTLs were not detected for SNPs in other candidate RNA editing genes.

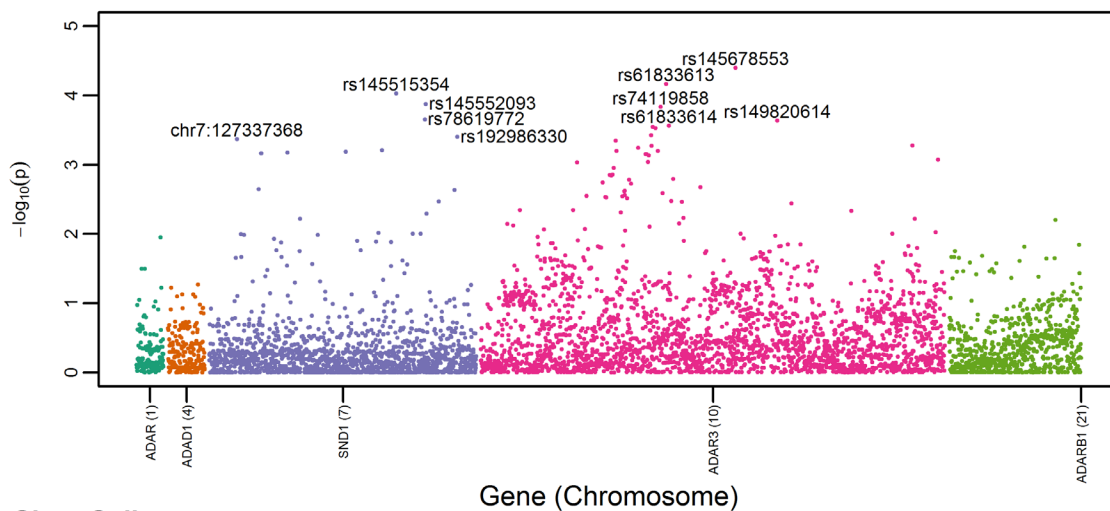
DISCUSSION

An emerging body of data suggest that defects in RNA editing may contribute to a range of human diseases, including cancer [2-4, 9-11]. The current large-scale collaboration represents the first comprehensive association study of germline variants involving RNA editing genes and susceptibility to epithelial ovarian cancer. At the SNP- level, the strongest associations were observed for SNPs in RNA editing genes *ADAR3* and *SND1*, but no associations reached genome-wide

A) Endometrioid



B) Mucinous



C) Clear Cell

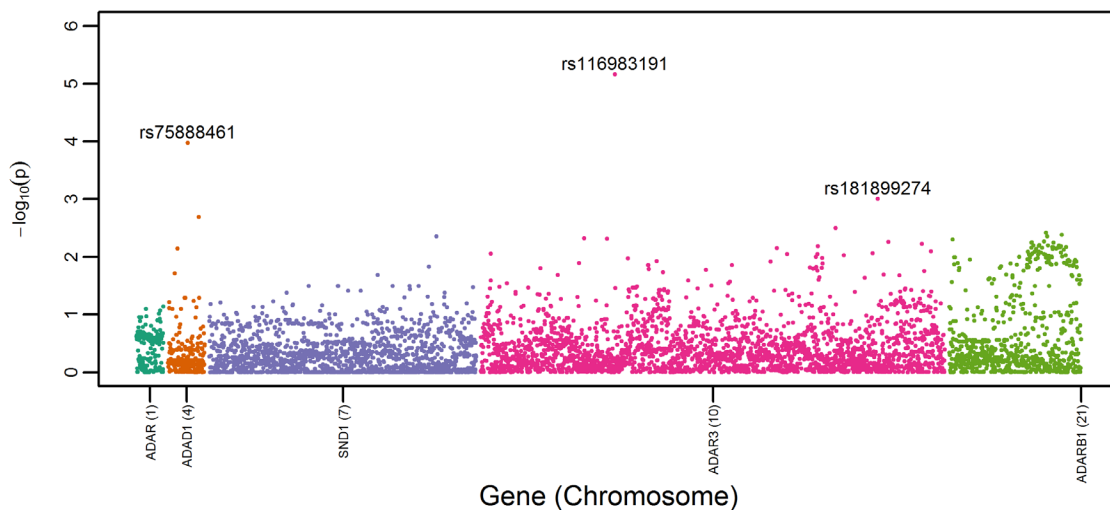


Figure 2: Manhattan plot for candidate RNA editing SNPs among a) endometrioid cases ($n = 1,439$) versus controls ($n = 21,693$), b) mucinous cases ($n = 696$) versus controls, and c) clear cell cases ($n = 660$) versus controls.

Table 2: Top-ranked RNA editing SNP-EOC risk associations among all histologies (N = 10,891) or serous histology (N = 6,500) versus controls (N = 21,693), sorted by gene and p-value

Gene	SNP	Alleles	MAF	Imputation accuracy R2	All histologies OR (95% CI)	P	FDR	Serous OR (95% CI)	P	FDR
ADAR	rs9426826	C>G	0.481	0.86	1.05 (1.02-1.09)	0.0038	0.76	1.04 (1-1.08)	0.0759	0.9996
	rs3738030	A>C	0.116	0.79	0.93 (0.89-0.98)	0.0080	0.83	0.91 (0.86-0.97)	0.0038	0.9996
ADAR3	rs77027562 ^a	A>G	0.009	0.41	1.39 (1.17-1.64)	0.0001	0.34	1.33 (1.08-1.62)	0.0061	0.9996
	rs11250601	C>T	0.070	0.62	0.89 (0.83-0.95)	0.0007	0.34	0.9 (0.83-0.97)	0.0071	0.9996
	rs142123280 ^b	A>G	0.001	0.48	2.08 (1.36-3.17)	0.0007	0.34	2.01 (1.23-3.29)	0.0056	0.9996
	rs4880912	T>C	0.200	0.82	1.07 (1.03-1.11)	0.0015	0.51	1.06 (1.01-1.11)	0.0267	0.9996
	rs11598359	C>T	0.005	0.45	0.68 (0.54-0.87)	0.0018	0.53	0.63 (0.47-0.86)	0.0032	0.9996
	rs6560760	C>T	0.025	0.59	1.17 (1.06-1.3)	0.0024	0.65	1.07 (0.95-1.22)	0.2688	0.9996
	rs2676202 ^c	C>T	0.122	0.66	0.93 (0.88-0.98)	0.0038	0.76	0.9 (0.85-0.96)	0.0014	0.9996
	rs139646191	TAGAA>T	0.062	0.66	1.11 (1.03-1.18)	0.0038	0.76	1.07 (0.98-1.16)	0.1242	0.9996
	rs139812582	G>A	0.002	0.47	1.71 (1.15-2.54)	0.0078	0.83	2.03 (1.31-3.14)	0.0017	0.9996
	chr10:1419524	T>TGG	0.009	0.60	0.79 (0.65-0.95)	0.0106	0.83	0.68 (0.54-0.86)	0.0014	0.9996
SND1	rs185147330	C>T	0.005	0.45	1.32 (1.05-1.67)	0.0176	0.86	1.54 (1.19-2)	0.0011	0.9996
	rs185455523	T>A	0.008	0.56	0.68 (0.56-0.83)	0.0002	0.34	0.6 (0.46-0.77)	0.0001	0.45
	rs145106202 ^d	G>C	0.009	0.87	0.73 (0.61-0.88)	0.0008	0.35	0.67 (0.53-0.84)	0.0006	0.9996
	rs181460088	C>T	0.008	0.80	0.73 (0.6-0.89)	0.0021	0.58	0.67 (0.52-0.86)	0.0020	0.9996
	rs199750392	G>GT	0.036	0.61	1.14 (1.05-1.25)	0.0028	0.71	1.11 (1-1.23)	0.0550	0.9996
	rs79138382	C>T	0.007	0.72	1.32 (1.09-1.61)	0.0056	0.83	1.42 (1.13-1.77)	0.0025	0.9996

Significant SNPs ($P < 4.0 \times 10^{-3}$) are listed and SNPs in LD ($r^2 > 0.60$) with more significant SNP are not reported:

- a) 7 SNPs in LD not reported
- b) 12 SNPs in LD not reported
- c) 1 SNP in LD not reported
- d) 1 SNP in LD not reported

levels of statistical significance. Gene-level analyses highlighted *ADAR* and *ADAR3* as potential contributors to EOC susceptibility within the set of ADAR-related genes. Finally, positive eQTLs were also observed between ADAR genotype and ADAR expression in EOC tumor tissue.

Focused evaluations of RNA editing SNP-disease associations are limited [12], especially with cancer as an outcome, so it is not possible to compare our SNP findings to those of other studies of cancer risk. We are, however, unaware of GWAS hits in or near these genes. Several recent studies [2, 3] have evaluated the genomic landscape and clinical relevance of RNA editing in numerous human tissue types. These analyses used RNA-sequencing data from both tumor and normal samples profiled as part of TCGA Project. Striking differences in RNA-editing patterns were observed in tumors relative to matched normal tissues for 12 cancer types [2]. Further analyses revealed that altered RNA editing patterns in tumors correlated with *ADAR* expression, and that non-random, clinically-relevant RNA editing events (frequently located in noncoding RNAs, nonsynonymous sites, intronic regions, and non-*Alu* elements) correlated with tumor classification and patient survival and with increased cell

survival and altered drug sensitivity [2, 3]. Interestingly, gene amplification-associated overexpression of *ADAR* was recently shown to enhance lung tumorigenesis and contribute to poor outcomes by affecting downstream RNA editing patterns [10]. As mentioned previously, ADAR expression levels have been reported to be significantly higher in serum and/or peritoneal fluid from patients with EOCs compared with benign ovarian tumors [6, 7]. Although high-grade serous EOCs from TCGA were not profiled as part of the aforementioned genomic investigations [18, 19], Haploreg 4.1 effectively integrates GTEx eQTL results for normal ovary.

Taken together with several lines of investigation from ovarian [6, 7] and other cancers [2, 3, 10] the current study suggests that ADARs (and *ADAR* in particular) may be useful biomarkers for the diagnosis and management of EOC. Thus, with replication, *ADAR* genotype status and/or expression level may serve as a risk factor for EOC. Indeed, we find that our top risk SNP in *ADAR*, rs9426826, has several proxy variants ($r^2 > 0.8$, Supplementary Table 2) that are strongly associated with expression of this gene in blood (rs1127313: 7.23×10^{-14}) and to a lesser extent, expression in high-grade serous EOC tumors (rs1127313: $P = 0.027$). Based on growing data which demonstrate

Table 3: HaploReg results for top-ranked ADAR3 SNP rs77027562 and its proxies from univariate analyses

Position (hg38)	LD (r ²)	SNP (Ref>Alt)	MAF in EUR	Functional Annotation	CR	Promoter histone marks	Enhancer histone marks	DNase site	Proteins bound	eQTL	Motifs Changed
Chr10:1688744	--	rs77027562 (A>G)	0.02	Intronic	No		BRN	BLD			ERalpha-a, RXRA, Zfp281
Chr10:1675149	0.94	rs12258319 (G>T)	0.98	Intronic	No		BLD				Pax-5
Chr10:1675875	0.94	rs7077743 (C>T)	0.98	Intronic	No		BLD				BDP1, CAC-binding-protein, HNF4, p300
Chr10:1676470	0.94	rs6560758 (T>C)	0.98	Intronic	No						
Chr10:1678882	0.94	rs10751814 (A>G)	0.98	Intronic	No						GR, Rad21
Chr10:1680294	0.94	rs7089727 (A>G)	0.98	Intronic	No		ESDR, IPSC, BLD, LNG	ESC, BLD	CTCF		
Chr10:1681695	0.94	rs6560759 (T>C)	0.98	Intronic	No		ESDR, BLD				Myc
Chr10:1687566	0.94	rs79784382 (T>A)	0.02	Intronic	No		BRN				Cphx, Duxl, HNF6, Hmx, Hoxa13, Pbx-1, Pbx3

Abbreviations: LD, linkage disequilibrium; MAF, minor allele frequency; EUR, European; CR, conserved region; eQTL, expression quantitative trait loci. Tissue groups: BRN, brain cells; BLD, blood and T-cells; ESDR, embryonic stem cell derived cells; IPSC, induced pluripotent stem cells; LNG, lung cell; ESC, embryonic stem cells. Proxies were defined as variants in LD ($r^2 > 0.8$) with the index SNP rs77027562 (bolded) in 1000 genomes project Phase 1 data for Europeans. All data was accessed using HaploReg v4.1 available at: http://www.broadinstitute.org/mammals/haploreg/documentation_v4.1.html. Both conservation prediction algorithms, GERP and SiPhy-omega, were used. Only eQTLs for ADAR genes (5 genes) are given.

Table 4: Association between RNA editing genes and EOC susceptibility.

Gene	Total N Markers (N Tested)	N Rare Markers (MAF<0.01)	N Common Markers (MAF≥0.01)	All Invasive		Serous		Endometrioid		Mucinous		Clear cell	
				P. SKAT-CR	P.AML Trend	P. SKAT-CR	P.AML Trend	P. SKAT-CR	P.AML Trend	P. SKAT-CR	P.AML Trend	P. SKAT-CR	P.AML Trend
ADAD1	210 (210)	98	112	0.698	0.749	0.300	0.433	0.054	0.179	0.857	0.804	0.696	0.635
ADAR1	155 (155)	50	105	0.020	0.022	0.110	0.101	0.005	0.008	0.943	0.841	0.780	0.411
ADAR2	754 (754)	301	4563	0.861	0.894	0.623	0.720	0.134	0.496	0.338	0.208	0.105	0.041
ADAR3	2656 (2654)	787	1867	0.216	0.266	0.334	0.541	0.587	0.470	0.002	0.031	0.234	0.502
SND1	1528 (1527)	764	763	0.630	0.809	0.703	0.376	0.919	0.895	0.632	0.204	0.773	0.535

the inhibition of tumor growth in the presence of ADAR inhibitors [13] and other therapeutic agents such as the IGFR-1R inhibitor BMS536924 and the MEK inhibitors CI1040 and trametinib [2], *ADAR* genotype and/or expression may help identify women whose tumors may respond to new combinations of therapies.

Strengths of the current study include the large sample size that primarily enabled detection of small effects for common variants, the relatively homogeneous population of EOC cases, and the multi-tiered genomic

evaluation. However, this study was underpowered to detect the rare variants that were identified and is burdened by the low imputation quality. Additionally, the study is limited in that eQTL analysis did not permit adjustment for somatic copy number changes and DNA methylation status, factors that can influence transcript abundance and confound associations between germline polymorphisms and gene expression [14-16]. Moreover, it is possible that the top-ranked SNPs could potentially affect genes other than the RNA editing genes that drive candidate

selection. Efforts to replicate these findings are needed; data will be available soon from a large, independent cohort of EOC cases genotyped by OCAC for this purpose (Amos et al, The OncoArray Consortium: a Network for Understanding the Genetic Architecture of Common Cancers (provisionally accepted, *CEBP*). Mechanistic studies to reveal how *ADAR* polymorphisms may affect oncogenic phenotypes will also be required, as will systematic investigations of the genomic landscape and clinical relevance of RNA editing in EOC using data from TCGA or other sources.

In summary, this study provides data to support the hypothesis that germline polymorphisms in *ADAR* related genes may influence gene expression and susceptibility to EOC. Further investigations are needed to determine whether inherited and acquired alterations affecting RNA editing serve as biological mechanisms to promote the development of EOC.

MATERIALS AND METHODS

Study population

A total of 41 studies (32 case-control and 9 case-only) from OCAC contributed to this investigation (Supplementary Table 1). Briefly, cases were women diagnosed with histologically confirmed primary invasive EOC (95%), fallopian tube cancer (1%), or primary peritoneal cancer (4%). Controls were women without

cancer and with at least one intact ovary on the reference date. Individual studies were grouped into 26 case-control strata. All studies provided data on disease status, age at diagnosis/interview, self-reported racial group, and histologic subtype.

Genotyping, quality control (QC), and imputation

Peripheral blood was the primary source of germline DNA and was collected in the course of clinical care or research at each of the participating sites. The candidate SNPs selected for the current investigation were genotyped using a custom Illumina Infinium iSelect Array as part of the international Collaborative Oncological Gene-environment Study (iCOGS), an effort to evaluate 211,155 genetic variants for association with cancer risk [17].

Briefly, OCAC genotyping was conducted at McGill University and Génome Québec Innovation Centre (Montréal, Canada) and Mayo Clinic Medical Genomics Facility. Each 96-well plate well contained 250ng genomic DNA (or 500 ng whole genome-amplified DNA). Raw intensity data files were sent to the COGS data coordination center at the University of Cambridge for genotype calling and QC using the GenCall algorithm. Sample and SNP quality control procedures have been described previously; in brief, samples were excluded with call rates < 95%, >1% discordance, < 80% European ancestry, or ambiguous gender, and SNPs were excluded with call rates < 95% or monomorphism [18, 19].

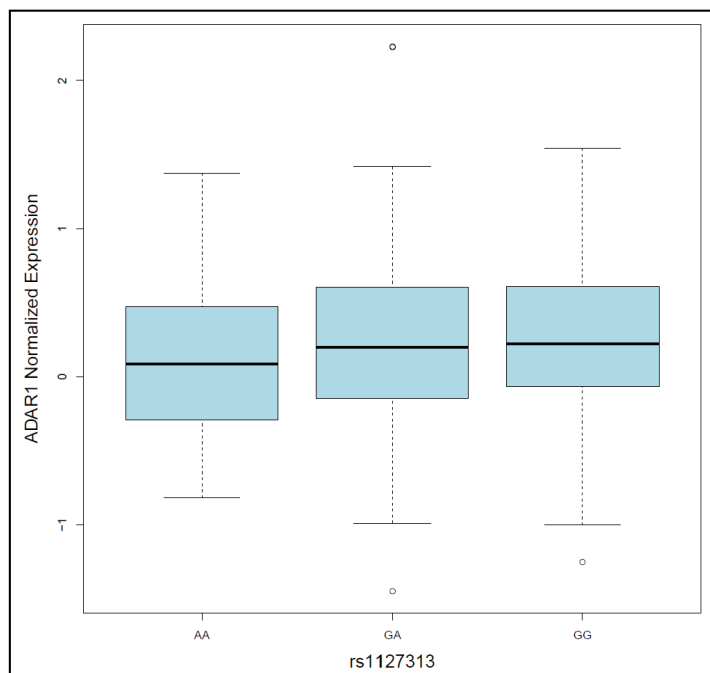


Figure 3: Box-plot showing that *ADAR1* tumor tissue expression differed, albeit only slightly, by rs1127313 genotype ($p = 0.027$).

To improve genomic coverage and power [14], we imputed genotypes based on data from the 1000 Genomes Project (1KGP); we used IMPUTE2 version 2 after pre-phasing with SHAPEIT [20]. All 14 populations in the 1KGP were used as the reference. Before imputation, we excluded poorly performing SNPs according to the genotyping success rates, deviation from Hardy-Weinberg equilibrium (HWE) ($P < 1 \times 10^{-7}$), and replicate errors. To ensure the quality of the imputed genotypes, maximum likelihood genotype imputation was carried out and an estimate of the squared correlation between the imputed and true genotypes was calculated. Imputation quality is significantly decreased for low and rare frequency variants [21]. To be more inclusive of rare variants, we considered imputed SNPs with an $r^2 > 0.40$ as well-imputed [22] and included them in our analyses. The average imputation quality for included variants is detailed in Supplementary Table 4, overall and by MAF categories.

Gene and SNP selection

Five candidate genes were chosen for this study based on published literature which directly showed or suggested roles in the regulation of A-to-I RNA editing [1, 4, 23]. The genes included adenosine deaminase domain containing 1 (*ADADI*), adenosine deaminase, RNA-specific (*ADAR/ADAR1*), adenosine deaminase, RNA-specific, B1 (*ADARB1/ADAR2*), adenosine deaminase, RNA-specific, B2 (*ADAR3/ADARB2*), and staphylococcal nuclease and Tudor domain containing 1 (*SND1*). In total, 5,303 SNPs in the 5 genes, 77 genotyped directly and 5,226 imputed, were available for statistical analysis.

Population stratification

HapMap DNA samples from European (CEU, $n = 60$), African (YRI, $n = 53$) and Asian (JPT+CHB, $n = 88$) populations were also genotyped as part of the same custom Illumina iSelect Array. The program LAMP [24] was used to estimate intercontinental ancestry based on the HapMap (release no. 23) genotype frequency data for these three populations. Eligible subjects with greater than 90 percent European ancestry were defined as European ($n = 39,773$). We then used a set of 37,000 unlinked autosomal markers to perform principal components analysis within each major population subgroup. To enable this analysis on very large sample sizes we used an in-house program written in C++ using the Intel MKL libraries for eigenvectors (available at <http://ccge.medschl.cam.ac.uk/software/>).

Statistical analysis

Descriptive statistics were calculated in terms of means and standard deviations for continuous variables and frequencies and percents for categorical variables. The primary association analysis focused on individuals of European ancestry. Unconditional logistic regression was used to estimate odds ratios (OR) and their 95% confidence intervals (CI) between genotype and case status under a log-additive genetic model, with adjustment for the first five principal components representing sub-European ancestry. Due to the heterogeneous nature of EOC, subgroup analyses were conducted to estimate genotype-specific odds ratios by histologic subtype: serous, endometrioid, mucinous, and clear cell carcinomas. False discovery rates (FDR) [25] were used to adjust for multiple comparisons, and FDR of 15% was used to declare significance.

Two methods of gene-level evaluations were also conducted to combine association evidence from SNPs within each gene evaluated: the Admixture Maximum Likelihood (AML) Test [26] and the Sequence-Kernel Association test for the combined effect of common and rare variants (SKAT-CR) [27]. AML is an approach that simultaneously examines the global null hypothesis (of no SNP-outcome associations) and estimates the proportion of underlying false hypotheses. The AML uses univariate SNP-level results to calculate the AML Cochran-Armitage Trend test. Compared to other methods, AML has been shown to have similar or higher statistical power to detect associations except under the unlikely scenario that greater than 20% of all variants are associated with the outcome [26]. SKAT-CR evaluates the cumulative effect of rare and common variants, but does not consider low-frequency variants. These gene-level approaches were undertaken to complement SNP-level findings, and aimed to reduce the degrees of freedom, avoid model-fitting issues due to multicollinearity from LD, and to improve statistical power. The Bonferroni method was used to account for multiple comparisons.

Expression quantitative trait locus (eQTL) analysis was performed to examine for association between genotype ($n = 5,303$, imputed as above in $n = 5$ genes) and corresponding gene expression for the 5 candidate RNA editing genes. Matched genotype and gene expression profiling data were obtained for 402 high-grade serous EOC samples evaluated in the Cancer Genome Atlas (TCGA) Project using previously described methods [19]. Briefly, germline genotypes and matched tumor gene expression data were downloaded from the TCGA data portal. To conduct the eQTL analysis, we used germline genotypes of SNPs/proxies as independent variables and expression levels as traits. Expression levels between minor allele carriers versus non-carriers were compared using the Wilcoxon rank sum statistic. Haploreg v4.1 <http://www.broadinstitute.org/mammals/haploreg/>

haplorege.php) [28] was used to evaluate the putative function of candidate SNPs.

CONFLICTS OF INTEREST

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